

**Method** A systematic review. We searched PubMed, EMBASE and Web of Science for the keywords “infectious diseases”, “research and development” and “pharmaceutical industry”.

**Preliminary results** The searches gave a total of 248 references. Among the findings, we want to highlight the Drugs for Neglected Diseases initiative (DNDi) and the WHO Research and Development Treaty (R&D Treaty). DNDi is a non-profit organization that has developed six new drugs since 2003. The development costs were €150 millions per drug, which is considerably below the costs for drug development claimed by the pharmaceutical industry. The R&D Treaty will commit member states of the WHO to fund development for neglected health needs using alternative incentives like milestone prizes, patent pools and direct grants. The treaty has not yet been agreed upon.

**Conclusions** Though a low priority from the pharmaceutical industry, other funding models have proven able to deliver new treatments. This could also lead to more development of non-patentable treatments, e.g. psychotherapy.

**Disclosure of interest** The author has not supplied his/her declaration of competing interest.

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#### EV1021

### Clinical and socio-demographic characteristics of a sample of outpatients with long-acting injectable antipsychotic treatment

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**Introduction** There are relatively few studies of Long-acting injectable antipsychotics (LAI), although poor adherence to treatment is one of the main problems in patients with psychotic disorders.

**Objectives** The aim of the study is to describe socio-demographic and clinical characteristics of a sample of outpatients with LAI treatment.

**Methods** This is a cross-sectional study. A randomized sampling was performed among the outpatients that were receiving LAI in an outpatient clinic in Barcelona (Spain). For each patient, socio-demographic, clinical and pharmacotherapeutic data were collected through interviews and clinical history.

**Results** The sample consisted of 30 subjects (50% men, average age 48 years). Most of the patients in the sample have basic education (50%) and are unemployed, receiving permanent disability pension (39.3%). In addition, 44.8% of the subjects were living with family members and were not married (56.7%). Of the patients, 70% were diagnosed with schizophrenia, 13.3% schizoaffective, 10% bipolar and 6.7% delusional disorder. The main reason to initiate LAI treatment was due to non-compliance of the prescribed oral treatment (85.7%). The 40% of patients were also with oral antipsychotic treatment. Average punctuation in the 3 first items of the Scale to Assess Unawareness of Mental Disorder: 11. Average punctuation in the short version of the Simpson-Angus Scale: 1.68.

**Conclusions** In our sample, the outpatients with LAI treatment had a low functioning and disease awareness. Although the main reason to start LAI is the non-compliance, 40% of the patients were concurrently treated with oral antipsychotics. The extrapyramidal side effects are mild.

**Disclosure of interest** The authors have not supplied their declaration of competing interest.

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#### EV1023

### Aripiprazole is effective for the improvement of psychotic symptoms in patients with dementia with lewy bodies

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**Objective** Dementia with lewy bodies (DLB) is commonly considered the second most common form of dementia. The purpose of this study is to investigate the treatment effects of aripiprazole in patients with DLB.

**Methods** Eleven patients who had meet the criteria for DLB participated in this study. The presence of psychotic symptoms was confirmed by scores of either the delusions or hallucinations items of the Neuropsychiatric Inventory (NPI) score. Patients who had 25 or more on the Mini-mental State Examination Scale (MMSE) at the entry or having brain damage were excluded. Aripiprazole was initiated at a low dose (3 or 6 mg/day) and titrated to higher doses at 2-weeks intervals or more rapidly based on investigator's judgment. Previous medications prior to aripiprazole administration were not changed through this trial. Patient's clinical status was assessed at baseline, then 2 weeks during the study by using NPI, Clinical Global Impression (CGI) and Brief Psychiatric Rating Scale (BPRS) to measure psychotic behavioral symptoms, and Simpson-Angus Scale (SAS) to measure parkinsonism symptoms. Clinical Dementia Rating (CDR) and MMSE were carried out at screening and end point to evaluate cognitive function.

**Results** The mean scores of the SAS and CDR were significantly decreased at the study endpoint compared to baseline. The mean scores of the NPI and BPRS improved up until 4 weeks after having started aripiprazole. After 4 weeks, improvements slowed. The mean score of the CGI-S was decreased up until 8 weeks.

**Conclusion** This study shows that aripiprazole may be effective for the treatment of psychotic symptoms in patients with DLB.

**Disclosure of interest** The authors have not supplied their declaration of competing interest.

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#### EV1024

### Tropicamide eye drops reduce clozapine-induced hypersalivation: A case report

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**Introduction** Clozapine-induced sialorrhea (CIS) is a common, treatment-limiting and stigmatizing side effect. All systemic agents that are used for hypersalivation may increase clozapine side effects such as blood pressure changes, constipation, or arrhythmias. Oral application of topical anti-muscarinic agents may be a low side effect option for treatment of CIS.

**Objective** The aim of this case report was to propose an off-label treatment of tropicamide drops to CIS and to stimulate further investigation.

**Case report** A 33-year-old male inpatient with schizophrenia has been on clozapine 800 mg and amisulpride 600 mg/day. His drooling was occasional and severe as drool drips off his chin during the day and night. Wet area over the pillow, visual analog scale (VAS), the short form of health survey (SF-36), UKU side effect rating scale, scale for the assessment of negative symptoms (SANS), scale for the assessment of positive symptoms (SAPS) were applied at baseline and in one-week intervals. Oral application of one drop of tropicamide % 0.5 (5 mg/mL) to left and one drop to right side